### PATENT COOPERATION TREATY PCT

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INTERNATIONAL PRELIMINARY EXAMINATION WEBORT (PCT Article 36 and Rule 70)

PCT

Applicant's or agent's file reference See Notification of Transmittal of International Preliminary FOR FURTHER M80516545:JWC:SRC:CT ACTION Examination Report (Form PCT/IPEA/416). International Application No. International Filing Date Priority Date (day/month/year) (day/month/year) PCT/AU2003/000771 20 June 2003 21 June 2002 International Patent Classification (IPC) or national classification and IPC Int. Cl. 7 A61K 31/7076, 31/167; A61P 41/00, 9/10; A01N 1/02 Applicant GLOBAL CARDIAC SOLUTIONS PTY LTD et al This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36. This REPORT consists of a total of 3 sheets, including this cover sheet. This report is also accompanied by ANNEXES, i.e., sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT). These annexes consist of a total of sheet(s). This report contains indications relating to the following items: I Basis of the report  $\mathbf{II}$ Priority Ш Non-establishment of opinion with regard to novelty, inventive step and industrial applicability ΙV Lack of unity of invention V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement VI Certain documents cited VII Certain defects in the international application VIII Certain observations on the international application Date of submission of the demand Date of completion of the report 20 January 2004 12 October 2004 Name and mailing address of the IPEA/AU Authorized Officer AUSTRALIAN PATENT OFFICE PO BOX 200, WODEN ACT 2606, AUSTRALIA E-mail address: pct@ipaustralia.gov.au Facsimile No. (02) 6285 3929 S. Chew

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I.	Basis of the report			
1.	With regard to the elements of the international application:*			
	the international application as originally filed.			
	the description, pages, as originally filed,			
	pages, filed with the demand,			
	pages, received on with the letter of			
	the claims, pages, as originally filed,			
	pages , as amended (together with any statement) under Article 19,			
	pages, filed with the demand,			
	pages, received on with the letter of			
	the drawings, pages, as originally filed,			
	pages, filed with the demand,			
	pages, received on with the letter of			
	the sequence listing part of the description:			
	pages , as originally filed			
	pages , filed with the demand			
	pages, received on with the letter of			
2.	ith regard to the language, all the elements marked above were available or furnished to this Authority in the language in nich the international application was filed, unless otherwise indicated under this item.  Lese elements were available or furnished to this Authority in the following language which is:			
	the language of a translation furnished for the purposes of international search (under Rule 23.1(b)).			
•	the language of publication of the international application (under Rule 48.3(b)).			
	the language of the translation furnished for the purposes of international preliminary examination (under Rules 55.2 and/or 55.3).			
3.	th regard to any <b>nucleotide and/or amino acid sequence</b> disclosed in the international application, the international oreliminary examination was carried out on the basis of the sequence listing:			
	contained in the international application in written form.			
	filed together with the international application in computer readable form.			
	furnished subsequently to this Authority in written form.			
	furnished subsequently to this Authority in computer readable form.			
	The statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.			
	The statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished			
4.	The amendments have resulted in the cancellation of:			
	the description, pages			
	the claims, Nos.			
	the drawings, sheets/fig.			
5.	This report has been established as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed, as indicated in the Supplemental Box (Rule 70.2(c)).**			
*	Replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to this report since they do not contain amendments (Rules 70.16 and 70.17).			
**	Any replacement sheet containing such amendments must be referred to under item 1 and annexed to this report			

## INTERNATIONAL PRELIMINARY EXAMINATION REPORT

nternational application No.
PCT/AU2003/000771

V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1.	Statement		
	Novelty (N)	Claims .	YES
	·	Claims 1-29	NO
	Inventive step (IS)	Claims	YES
	Industrial applicability (IA)	Claims 1-29 Claims 1-29	NO
		Claims 1-29	YES
_	0:		NO

2. Citations and explanations (Rule 70.7)

This report has considered the following documents cited in the International Search Report:

- D1 WO 2000/056145 A
- D2 Derwent Abstract Accession Number 74319 E/35
- D3 US 5407793 A

#### NOVELTY (N): Claims 1-29

D1 has disclosed a composition and its use thereof for arresting, protecting or/and preserving an organ (eg heart) comprising (1) a potassium channel opener or an adenosine receptor agonist, (2) a local anaesthetic and (3) a suitable agent such as sucrose, vitamin E, alpha - tocopherol, ascorbic acid and magnesium stearate (See the examples, claims and page 13 line 4-20).

Therefore claims 1-29 lack novelty.

D2 has disclosed a heart perfusion aqueous composition useful for keeping the isolated heart functioning comprising adenosine triphosphoric acid, lidocaine, glucose, mannitol, heparin, magnesium asparginate, succinate and bicarbonate salts.

D3 has disclosed an aqueous heart preservation solution which includes adenosine, lidocaine, glucose, histidine, calcium, sodium and magnesium salts (See examples).

Therefore claims 1-6 and 15-29 lack novelty in the light of the disclosures of D2 and D3 respectively.

## INVENTIVE STEP (IS): Claims 1-29

As above

# INDUSTRIAL APPLICABILITY (IA): Claims 1-29

Claims 1-29 have industrial applicability.